

Pregnant and non-pregnant women and low back pain-related differences on postural control measures during different balance tasks

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ABSTRACT

Introduction: Low back pain (LBP) is the most common musculoskeletal complaint in pregnancy, being responsible for many negative impacts. **Objective:** To evaluate the effect of LBP on static and dynamic balance in pregnant women and whether pregnancy mediates the results compared to non-pregnant women. **Methods:** 44 women (mean age 30 yrs) participated voluntarily in this study: 16 pregnant women with LBP starting in pregnancy, 14 pregnant women without LBP and 14 non-pregnant women as a group control. Participants were assessed for static postural balance using a force platform and dynamic mobility balance using the Timed Up and Go (TUG) test. **Results:** The pregnant women with LBP showed significant ($P < 0.04$, for mean, $d = 1,2$) poor postural balance in static tests (force platform), in the area of COP eyes open. In dynamic balance (TUG test), statistical difference was found between the groups ($P 0.038$) and the effect size were moderate to strong in the comparison between the three groups. The most sensitive differences were reported mainly between pregnant women with LBP versus non-pregnant control group in balance measures from force platform. **Conclusion:** The findings indicate that LBP associated to pregnant clinical status can decrease the balance capacity in women. These results have implication for balance evaluation and retraining in pregnant women with and without LBP from rehabilitation or prevention programs.

Keywords: Low Back Pain; Pregnancy; Postural Balance; Pain; Posture.

BACKGROUND

Low back pain (LBP) is one of the most frequent cause of disability affecting the worldwide adult population⁽¹⁾. Clinical observations suggest that approximately 84% of adults will experience a LBP at some point in their life, in Brazil, each year, it was estimated that up to 65% of men and women in all ethnic groups suffered from LBP⁽²⁾. LBP is defined as specific or non-specific axial or sagittal musculoskeletal pain located between L1 to L5 vertebrae with or without radiating leg symptom⁽³⁾. Pain severity correlates with function, those with more severe pain had poorer function while those with mild, well-controlled pain functioned normally⁽⁴⁾.

LBP genesis may have diverse origins, such as repetitive movements, postures, stress or exposition of prolonged static position, mechanical, hormonal, circulatory and psychosocial issues which are now well-known factors capable of increasing the risk of chronic pain⁽⁵⁾. Based on that fact, it is no surprise that the specific LBP estimated incidence differs during pregnancy. In fact, pregnancy-related LBP is a common symptom affecting 24% to 90% of these women worldwide. Because of the lack of knowledge and understanding of LBP, health care professionals do not know how to deal with or

relieve patient's symptoms. LBP are often sufficient to alter normal physical and daily life activities^(6,7).

During the pregnancy period, many biological adaptations such as skeletal muscle mass, body dimensions and posture change the morphological characteristics of women and in turn, may increase spinal instability and generate back pain⁽⁸⁾. From a biomechanical perspective, these adaptations move the center of gravity (COG) to the anterior and superior poles in body and may contribute to balance disorders. In fact, static balance is the ability to keep the COG within the base of support and dynamic balance as the capacity to perform a task while restoring a stable COG position⁽⁹⁾. Under normal condition, balance performance is maintained by both static and dynamic controls. The pregnancy-related COG displacement can affect postural control strategies related to neuromuscular changes and changes in biomechanics for equilibrium of the body⁽¹⁰⁾. Apparently, postural alterations are evident during the gestational period, as well as expected changes in balance⁽¹⁰⁾. However, this phenomenon increases the risk of musculoskeletal discomfort in the thorax and lower limbs and consequently, causes foot, back and lower limb disorders, gait changes, poor

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mobility and pain⁽¹¹⁾. Therefore, the repercussions of pregnancy on the musculoskeletal system result in great adjustments in static and dynamic posture for a woman⁽¹²⁾.

Few studies have, however, compared balance and mobility in pregnant women with LBP and these results relative to a control non-pregnant group⁽¹¹⁾. Opala-Berdzik et al. investigated static stability measures under different conditions in women between the onset of gestation, advanced gestation and 2 and 6 months post-gestation. These authors reported a significant negative balance effect in the advanced gestation group when compared to the non-gestational state⁽⁸⁾. It would be interesting to compare these results with pregnant women with LBP as well as with non-pregnant women.

The aim of this study was to compare static and dynamic balance in pregnant women with and without LBP and non-pregnant women. The hypothesis of this study was that pregnant women with LBP would present poorer functionality and postural balance than women without LBP or women not pregnant.

METHODS

Design and participants

This research was an exploratory and descriptive cross-sectional study with a convenience sample. The study included pregnant and non-pregnant women. For pregnant women with LBP, the inclusion criteria were: 1) women over 18 years old; 2) between 22 and 33 weeks of gestation; 3) being under prenatal clinical follow-up; 4) nonspecific LBP reported with Visual Analogic Scale (VAS) 3 to 10; 5) not having or participating in specific treatment (physical therapy or medication for LBP in the last 3 months); 6) able to perform functional activities, and not presenting limitations in relation to cognition and attention. For pregnant women without LBP, the inclusion criteria were: 1) women over 18 years old; 2) between 22 and 33 weeks of gestation; 3) being under prenatal clinical follow-up; 4) nonspecific LBP reported with VAS 0 to 2; 5) not having or participating in specific treatment (physical therapy or medication for LBP in the last 3 months); 6) able to perform functional activities, and not presenting limitations in relation to cognition and attention. For non-pregnant women, were: 1) women over 18 years old; 2) not having gestation in the last 12 months; 3) without or not reporting LBP in the past 3 months; 4) able to perform functional activities and not presenting limitations in relation to cognition and attention. For both cases (pregnant and non-pregnant women) the exclusion criteria were: inability to perform the tests proposed or showing any condition that indicated a risk pregnancy. The study and measurement protocols were approved by the local ethics committee (number 1.579.189).

Based on a previous balance study with pregnant women using the center of pressure (COP) variable of the force platform: Velocity anteroposterior, pregnant women = 5.36 ± 0.25 mm/s and in non-pregnant women Velocity anteroposterior = 4.96 ± 0.23 mm/s, the minimal

sample size for a power of 0.95 at the 0.05 significance level was 11 participants per group⁽¹³⁾. A total of 30 pregnant and 14 non-pregnant women participated in this study. The pregnant women were divided into two groups according to VAS score for LBP, pregnant women with LBP (G1) for score between 3 and 10 ($n=16$, age in mean 32; Standard Deviation: SD = 7 yrs), pregnant women without LBP (G2) for score between 0 and 2 ($n=14$, age in mean 29; SD = 6 yrs), and non-pregnant women without LBP (G3), as a control group ($n=14$, age in mean 30; SD = 7 yrs).

Procedures and experimental protocol

For the evaluation of participants, pregnant women, a standardized protocol was used for data collection including personal information, anthropometric measures, VAS, and obstetric history, as well as evaluation of the presence of pain through physical examination and tests of the mobility of the spine and pelvis⁽¹⁴⁾. For pregnant women were still used, three validated questionnaires were used to assess the clinical state: 1) disability related to pain using the Roland-Morris Disability Questionnaire (RMDQ)⁽¹⁵⁾ and the Oswestry Disability Index (ODI)⁽¹⁶⁾ and 2) the pain state using the McGill Pain Questionnaire (MPQ)⁽¹⁷⁾. To ensure a healthy control group (non-pregnant women), VAS, personal data and anthropometric measurements were collected. All participants in the control group reported zero on the VAS measure.

Static balance assessment

For static balance (Figure 1), we used postural control measures from a force platform (BIOMECH 400, EMG system do Brasil, SP, Ltda). Two static balance experimental task conditions were performed randomly: (1) two-legged stance with eyes



Figure 1. Illustration of a typical participant under force platform for balance measurement (BIOMECH 400). Note: Static balance experimental task conditions: two-legged stance with eyes open and with eyes closed. 50x42mm (300x300 DPI).



open (TLEO); (2) two-legged stance with eyes closed (TLEC). All participants were familiarized with the equipment and balance protocol until they were comfortable with the testing. Balance assessment was performed with a standardized protocol: barefoot with arms at the sides or parallel to the trunk. During testing with eyes open, the participant would look at a target (a cross) placed on a wall at eye level 2 m away. To prevent falls during testing, an investigator stood close to the volunteers during all tasks. For each balance condition, three trials of 30 s with 30 s rest intervals were performed and the mean was retained for analysis⁽¹⁸⁾. A landmark on the force platform was used to standardize the position of the feet during all balance conditions.

The vertical ground reaction force data from the force platform was sampled at 100 Hz. All force signals were filtered with a 35- Hz low-pass second-order Butterworth filter. The signals from four sensors were converted into COP data using computerized stabilography, which was compiled with MATLAB routines (The Mathworks, Natick, MA). Stabilographic analysis of COP data led to the computation of the main balance parameters: the 95% confidence ellipse area of COP (A-COP in cm²), mean velocity (VEL in cm/s) in both anteroposterior (A/P) and mediolateral (M/L) directions of movement, and total displacement of COP (cm)⁽¹⁹⁾.

Dynamic mobility balance assessment

For physical mobility tests, we used the Timed Up and Go (TUG) test, which was undertaken using a chair with seat height of 46 cm, back-support and armrests. The participants were oriented “to walk as fast as you can until you cross the cone, turn around, and walk back to the chair and sit down again”. The cone was positioned 3m from the patient’s starting position. The time taken to complete the TUG was recorded in seconds using a chronometer bio Stopwatch (Model No. SW 2018). A total of two trials were assessed with 1 minute intervals between each trial, the best time performance being considered as a final measure⁽²⁰⁾. A trained physiotherapist assessed the postural balance and TUG tests.

STATISTICAL ANALYSES

Categorical variables were described in absolute frequency (n) and relative frequency (%), while numerical variables were described in mean and standard deviation (SD). Shapiro-Wilk test was used to evaluate the normality of the balance variables and determine which tests would be used. One-way ANOVA was used to assess differences between groups (pregnant women with LBP x pregnant women without LBP x control) related to balance measures (force platform and TUG). The effect size (*d*) between groups was calculated to determine the magnitude of effects using the equation: $d = m1 - m2 / SDm2$, where *m1* is the mean of the pregnant women with LBP group, while *m2* is the mean of the control group, and *SDm2* is the standard deviation of the control

group. The effect size was characterized as by Cohen (2013) as weak, moderate and strong effects, i.e., *d* = 0.2 is small, *d* = 0.5 medium and *d* = 0.8 large, respectively⁽²¹⁾. Spearman Correlation analysis was applied between the results of the questionnaires and the data of the force platform and TUG. All statistical analyses were performed with SPSS 20.0 for Windows (SPSS Inc., Chicago, IL, USA) with a level of significance of 0.05.

RESULTS

The three groups were homogeneous for anthropometric characteristics (age, weight, height and body mass index), with no significant difference (*P* > 0.05) between them. Table 1 also shows the data related to parity and pain characteristics of women. The descriptive results (table 1) showed that 31% of G1 pregnant had a score that indicates disability by RMDQ. In the ODI questionnaires, 50% of G1 pregnant obtained scores 21 to 40, indicating a moderate disability. The VAS pain score (0-10) for G1 pregnant had mean of 6, and for G2 pregnant was 1.5. In the questionnaire (MPQ) the most frequent type of pain for G1 pregnant was described “tiring / exhaustive” (76%).

Pregnant women of G1 and G2 (pregnant) showed poor static and dynamic balance, for all variables analyzed when compared to G3 (no pregnant). The most significant and sensitive differences between groups were reported for the force platform measurement, in Area of COP eyes open, where *G1* > *G3* *d* = 1,2 and *G2* > *G3* *d* = 0.90 (*P* 0.040); while for the eyes closed, was no statistically significant difference, but the effect size was moderate to strong across these differences (*G1* > *G3* *d* = 0.83 and *G2* > *G3* *d* = 0.72); in the other variables of the force platform there was no statistically significant difference between the groups (table 2) and the effect size was weak to moderate (*G1* x *G3* *d* = 0.7 to 0.50; *G2* x *G3* *d* = 0.23 to 0.53; *G1* x *G2* *d* = 0.1 to 0.27). For the TUG, statistical difference was found between the groups (*P* 0.038) and the effect size were moderate to strong in the comparison between the three groups (*G1* x *G3* *d* = 0.86; *G2* x *G3* *d* = 0.53; *G1* x *G2* *d* = 0.54).

The results of correlations of clinical status (pain and disability scores) and COP parameters and TUG varied from weak to moderate across the groups, where among the questionnaires and variables of the platform *r* was .36 to .43, between questionnaires and TUG *r* was .09 to .22, and between platform and TUG *r* was .00 to .26.

DISCUSSION

This study aimed to compare the static and dynamic postural balance of pregnant women with and without LBP compared to non-pregnant women. LBP during pregnancy is one of the most common musculoskeletal condition. Similarly than epidemiological data, in the present investigation, 53% of pregnant women experience significant LBP (VAS ≥ 3). In addition, pregnant women with and without LBP have poor postural balance when compared to non-pregnant women,

**Table 1.** Characterization of pregnant and non-pregnant women.

Characteristic	Pregnant	Pregnant	No pregnant woman
	G1	G2	G3
	(16)	(14)	(14)
Age (years)	32 (7)	29 (6)	30 (7)
BMI (kg/m ²)	25 (2)	25 (3)	24 (5)
VAS	6 (1)	1.5 (1)	0
Gestational age (weeks)	23 (5)	23 (5)	-
Uterine fundus (cm)	40 (5)	38 (6)	-
Abdominal circumference (cm)	91 (11)	93 (8)	-
Lumbar pain time (months)	3 (2)	2 (4)	-
Primiparous	(5) 31%	(12) 85%	-
Occupation			-
Housewife	(6) 37%	(2) 14%	-
Teacher	(3) 19%	(4) 28%	-
Student	(1) 6%	(2) 20%	-
Other occupation	(6) 37%	(6) 43%	-
Radiated pain reported	(7) 48%	(2) 14%	-
Activity practice before pregnancy	(7) 48%	(7) 50%	-
Have pain in rest	(13) 81%	(4) 29%	-
Have night pain	(5) 31%	(3) 21%	-
What makes the pain worst?			-
Seated position	(2) 12%	(4) 29%	-
Standing position	(2) 12%	(2) 14%	-
Maintain position for an extended period	(4) 25%	(1) 7%	-
Others	(8) 50%	(1) 7%	-
What makes the pain better?			-
Rest	(7) 48%	(6) 43%	-
Change position	(3) 19%	(1) 7%	-
Others	(6) 37%	(1) 7%	-
Roland Morris Questionnaire			-
From 0 to 6 points	(3) 19%	(7) 50%	-
From 7 to 13 points	(8) 50%	(4) 29%	-
From 14 or more points	(5) 31%	(3) 21%	-
Oswestry Questionnaire			-
From 0 to 20%	(5) 31%	(9) 64%	-
From 21 to 40%	(8) 50%	(4) 29%	-
From 41 to 60%	(3) 19%	(2) 14%	-

Note: Low Back Pain (LBP); Body Mass Index (BMI); Visual Analogue Scale (VAS). Mean (Standard deviation) of the characterization pregnant women with LBP (G1), without LBP (G2) and no pregnant and without pain (G3), in relation to anthropometric and personal data, percentage of characterization of pain, and pain and disability scores.

and the results for the effect size for these differences, indicating that LBP is a factor for decreasing balance in pregnant women. To the authors' knowledge, there are few studies that to compare three different groups of women in the same experimental design, which supports the originality of the results found. Past studies addressed this question

separately, analyzing different gestational periods⁽⁵⁾ or LBP effects post-childbirth^(22,23)

In brief, the characteristics of our sample (Table 1) were similar to the other studies related to LBP in pregnancy^(5,6,24), as well as the factors that interfere in the improvement or worsening of pain⁽²⁵⁾. The LBP women group had a moderate

**Tabela 2.** Mean (Standard deviation) for timed up go test and postural balance variables.

Postural balance		Pregnant	Pregnant	No pregnant	Anova	Tukey
		G1	G2	G3	F (P value)	(direction)
A-COP (cm ²)	EO	2.34 (1.54)	2.94 (2.80)	1.11 (0.64)	3.48 (0.04)*	<0.05 (Pregnant G1 > Control)
	EC	4.28 (5.1)	2.87 (3.34)	1.15 (0.55)	2.78 (0.73)	<0.05 (Pregnant G1 > Control)
VEL A/P (cm/s)	EO	0.73 (0.21)	0.76 (0.20)	0.71 (0.13)	0.32 (0.72)	--
	EC	0.92 (0.21)	0.93 (0.25)	0.83 (0.16)	1.01 (0.37)	--
VEL M/L (cm/s)	EO	0.55 (0.07)	0.53 (0.08)	0.51 (0.09)	0.72 (0.49)	--
	EC	0.59 (0.07)	0.63 (0.29)	0.53 (0.17)	1.13 (0.33)	--
Total-D (cm)	EO	61.84 (10.85)	61.2 (13.54)	57.23 (9.18)	0,70 (0.50)	--
	EC	72.03 (11.98)	71.59 (15.54)	64.69 (10.11)	1.50 (0.23)	--
TUG (s)		7.51 (1.93)	6.67 (0.92)	6.20 (0.85)	3.55 (0.03)*	<0.05 (Pregnant G1 > Control)

Note: Timed up and go test (TUG); Area of center of pressure (A-COP); COP velocity in the anteroposterior (VEL A/P) and mediolateral (VEL M/L) directions; Total displacement (Total-D); Low back pain (LBP); Eyes open (EO); Eyes closed (EC). *Significantly different.

disability and pain intensity level (Table 1). However, correlations between clinical status (disability and pain questionnaires) and measures of balance were poor to moderate. These results confirm that changes in balance measures may be related to a combined effect of pregnancy itself and the pathology of LBP. Moreira et al, evaluated 15 pregnant women with LBP and 15 non-pregnant women, concluded that the changes in the postural oscillation of the pregnant woman may be related to changes in biomechanical and hormonal levels, in addition, he further concluded that postural control during static posture can't be used to predict the occurrence of LBP⁽¹³⁾.

Our results for balance are in agreement with some studies^(9,18,19) and in disagreement with others^(10,12,23). For example, Opala-Berdzik et al evaluated 31 pregnant women without LBP and reported no effect on postural balance with eyes open. These authors reported an effect of an increase of postural instability only with the eyes closed and during advanced pregnancy (around the 36th week), which is contrary to the results of the present study, which found a decrease in both open and closed eyes, and also found that this balance decreases as LBP is higher⁽²⁴⁾.

We observed to a statistical difference in the TUG test when comparing pregnant women with LBP and non-pregnant women (Table 2), our results showed that pregnancy itself worsens mobility, but the presence of LBP has impact on the decrease of mobility during pregnancy. This is in agreement with a previous study which showed that difficulties in sitting and standing are significant in pregnancy and that these conditions may worsen due to LBP⁽⁶⁾. All together, these results suggest that effects on balance can be task dependent. More challenging balance performance and sophisticated equipment are suitable to better discriminate differences between pregnant women with LBP and non-pregnant women, as the present study did. In this sense, the strength of our experimental protocol was the use of a clinical protocol

related to direct postural control measurements through a stabilographic analysis on a force platform, which determined the difference in balance across groups.

On the other hand, the causes of LBP are not fully understood, but during pregnancy, we must consider a series of factors related to hormonal and biomechanical changes⁽²⁵⁾, as well as other factors related to increase in body mass index, decreased muscle strength and center of gravity changes^(22,23). According to Vermani et al., an increase of movement from pelvic joints in pregnant women, decreases the efficiency of load transmission, increases shearing forces between pelvic joints and leads to a higher probability of pain⁽²⁶⁾. In fact, one explanation for poor balance in pregnant women with LBP can be associated with these factors.

In addition, Panjabi's model⁽²⁷⁾ can also explain at least some of our results. Some studies of people with LBP have reported significantly poorer balance across this population when compared with health controls⁽¹⁹⁾. The differences between the groups with and without LBP may be related to trunk muscle fatigue in individuals with LBP, resulting in brief uncontrolled intervertebral movements and postural instability⁽²⁸⁾. Individuals with LBP present low levels of trunk muscle resistance compared to controls⁽²⁹⁾. In addition, impaired lumbosacral proprioception is also associated with LBP⁽³⁰⁾, which in turn affects balance performance.

We found a large effect size for the differences between groups of pregnant women with LBP and without LBP, compared to control, but we should note that the effect size was higher when we analyzed the group with LBP compared to the control, which indicates that pregnancy itself is a sufficient factor for decreasing the balance, and the presence of LBP increases this instability. However, currently there is no normative data for what is the minimal clinically important difference (MCID) in balance based on center of pressure measures. This is an important topic and future studies are needed to address this scientific gap.



Finally, the clinical implications of our results is a greater understanding of LBP characteristics during pregnancy and its repercussions, and thus can guide the professional in the formulation of physiotherapeutic objectives and activities that address the set of symptoms and deficits that pregnant women with LBP experience. Another implication is to demonstrate that pregnancy associated with LBP is a determinant factor in the decrease in functionality and the presence of an incapability level as shown in the results from the applied questionnaires. Thus, physiotherapists could better address balance evaluation and retraining for pregnant women with LBP in prevention or rehabilitation programs.

One limitation of this study was to find pregnant women totally without LBP. After a more careful evaluation, even with the absence of self-reported pain, the anamnesis and the physical examination detected the presence of a minimal level, so the study accepted the level of VAS \leq 2, as absence of pain. Other equilibrium tests (eg, semi-tandem) were not evaluated. Therefore, the results may not be generalizable for the entire population. However, we combined participants in significant demographic variables to reduce potential confounding factors, but other variables not used in the match may still have affected comparability between groups.

CONCLUSION

Pregnant women with and without LBP presented poor postural control and lower mobility when compared to non-pregnant women, where the presence of LBP increased these differences. These results have implications for the assessment of balance and retraining in pregnant women with and without LBP from rehabilitation or prevention programs.

AUTHOR'S CONTRIBUTIONS

APFC conceived of the study, and participated in its design and coordination, and drafted the manuscript; MRO participated in the design of the study and performed the statistical analysis, and helped to draft the manuscript; FKS, PEAS and RSS participated in the collection of evaluation data and experimental protocol; SSD, MD, MD and SN participated in its design and helped to draft the manuscript; RAS conceived of the study, and participated in its design and coordination, performed the statistical analysis and drafted the manuscript. All authors read and approved the final manuscript.

CONFLICT OF INTEREST

The authors declare that they have no competing interests.

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